

## PHARMACEUTICAL COMPOSITION OF AMLODIPINE MALEATE HAVING ENHANCED STABILITY

### Technical Field

5           The present invention relates to formulation-stability of amlodipine maleate. The present invention may provide a coated particle of amlodipine maleate and pharmaceutical composition thereof with formulation-stability equal to or higher than amlodipine besylate regardless of pH of composition during long storage due to coating a particle of amlodipine maleate with pharmaceutically acceptable coating  
10   agent and preventing decomposition reaction of amlodipine maleate.

### Background Art

          It is known that decomposition reaction of drug is complexly affected by pH, water or crystalline phase, purity or particle size of drug and the like.

15           Amlodipine is useful compound as calcium channel blocker for treating a wide variety of cardiac conditions, particularly cardiovascular disease (CVD) including angina, hypertension, etc.

20           European Patent Laid Open Publication Number 89,167 has disclosed various forms of pharmaceutically acceptable salts of amlodipine, particularly pharmaceutically acceptable acid addition salts, such as hydrochloride, hydrobromide, sulfate, phosphate or acid phosphate, acetate, maleate, fumarate, lactate, tartrate, citrate or gluconate, preferably maleate.

25           However, formulations comprising amlodipine maleate are not commercially developed due to poor stability by decomposition reaction during storage.

          Korean Patent Laid-Open Publication Number 1995-7228 has disclosed

amlodipine besylate that may overcome problems related with formulation-stability of amlodipine maleate.

In the prior art, stability on besylate, mesylate, succinate, salicylate, maleate, 5 tosylate, acetate and hydrochloride of amlodipine was evaluated using a mixture, as vehicle of tablet, of microcrystalline cellulose of 50:50 and anhydrous dibasic calcium phosphate, and as a result, it is disclosed that amlodipine besylate is most stable.

Drug comprising amlodipine has been approved as amlodipine besylate. 10 Although Pitcher co., Ltd. initially had applied amlodipine maleate to United States Food and Drug Administration (US FDA), converted amlodipine maleate from amlodipine besylate when US FDA reviewed effective ingredient of the drug.

However, amlodipine besylate has problems that bioavailability of each individual represents in a large range of about 64 to about 90 % due to high 15 deviations of solubility and dissolution rate under pH conditions. And benzene sulfonate, with corrosive wear on skin or eyes and respiratory toxicity due to inhalation, is used during manufacture of amlodipine besylate, thereby to cause stability problem on workers and environmental toxicity during synthesis of drug.

20 To enhance formulation-stability of amlodipine, development of new salts of amlodipine has been tried. Korean Patent Laid-Open Publication Number 2002-76561 has disclosed amlodipine camsylate manufactured by reaction with amlodipine and camphorsulfonic acid, Korean Patent Laid-Open Publication Number 2003-81006 has disclosed amlodipine nicotinate, Korean Patent Laid-Open 25 Publication Number 2004-11752 has disclosed tetrahydro-5-oxo-2-furane carboxylate of amlodipine, Korean Patent Laid-Open Publication Number 2003-17380 has disclosed a pyroglutamic acid salt of amlodipine, Korean Patent Laid-

Open Publication Number 2004-72363 has disclosed a cyclamate of amlodipine. However, the new salts were developed related with the patent for amlodipine besylate and the try is weak to solve the problem for formulation-stability of amlodipine maleate evaluated the most preferred salts.

5 Further, Korean Patent Laid-Open Publication Number 2003-70594 has disclosed that a pharmaceutical composition, comprising amlodipine maleate, has excellent stability at a pH of about 5.5 to about 7.0, when measured as aqueous slurry of 20 weight percent.

10 Although the pharmaceutical composition comprising amlodipine maleate has efficiencies on stability due to selecting excipients so as to having of the pharmaceutical composition at a pH of about 5.5 to about 7.0, the stability of the pharmaceutical composition has equal to or less than one of amlodipine besylate according to test of stability described in Korean Patent Laid-Open Publication Number 2003-70594.

15 Korean Patent Publication Number 1995-7228 has disclosed that amlodipine maleate is unstable than amlodipine besylate, based on evaluation of stability of a pharmaceutical composition comprising amlodipine maleate and a pharmaceutical composition including amlodipine besylate when pH of the pharmaceutical compositions has a range of about 5.5 to about 7.0.

20 The present inventors has discovered that amlodipine maleate may be easily decomposed by direct contact with excipients during studying on amlodipine maleate with improving bioavailability due to high solubility. Accordingly, the present inventors prepared amlodipine maleate by coating to thereby preventing  
25 decomposition reaction of amlodipine maleate without direct contact of excipients and/or regardless pH of a pharmaceutical composition comprising amlodipine maleate.

## Disclosure of the Invention

### Technical problem

To overcome problems of amlodipine maleate, that has excellent  
5 bioavailability due to high solubility thereof but commercially not developed due to  
decomposition reaction of drugs containing amlodipine maleate, it is an object of the  
present invention to provide a pharmaceutical composition with formulation-stability,  
comprising amlodipine maleate, for treating cardiovascular disease (CVD).

### 10 Technical solution

To accomplish the above-mentioned object, the present invention provides a  
coated particle of amlodipine maleate.

The present invention also provides pharmaceutical composition, comprising  
the coated particle of amlodipine maleate as active substance and pharmaceutically  
15 acceptable carriers, for treating cardiovascular disease (CVD).

The pharmaceutical composition comprising the coated particle of  
amlodipine has excellent bioavailability and may be sold commercially due to  
improving formulation-stability during long storage regardless pH of the  
20 pharmaceutical composition. Preventing decomposition reaction of amlodipine  
maleate and decreasing a loss of amlodipine equal to or more than a loss of  
amlodipine besylate cause improvement of formulation-stability.

Formulations containing amlodipine are prepared for rapidity and stability  
after oral dosing such that formulations need to have sufficient dissolution rate in  
25 vitro. Therefore the pharmaceutical composition comprising the coated particle of  
amlodipine maleate of the present invention further is necessary to be prepared for  
coating of amlodipine maleate without decreasing a dissolution rate in vitro.

The pharmaceutical composition comprises the coated particle with a dissolution rate of about 80 weight percent by a dissolution experiment, which is carried to using 0.01mol/L hydrochloric acid aqueous solution of 500mL for about 30minutes at about 75 rpm by conventional methods.

5        The pharmaceutical composition comprising the coated particle of amlodipine maleate has excellent bioavailablility as well as formulation-stability equal to or more than a pharmaceutical composition comprising amlodipine besylate.

10        In part of pharmacy, drugs is conventionally coated using coating agent for intercepting unsavory and a bad smell, enhancing a value of product by change of color, controlling dissolution amount of drugs, and protecting acid-laid drugs from acid in the stomach for oral-dosing and unstable drug from exterior circumstances such as water, light, etc. And the drugs are coated as a unit formulation, such as granule, tablet, etc.

15        Although a pharmaceutical composition comprising amlodipine maleate is coated as a unit formulation by a conventional coating method, amlodipine maleate is decomposed. In Korean Patent Laid-Open No. 2003-70594, is disclosed a pharmaceutical composition comprising amlodipine maleate and excipients selected properly in order to have a pH range of about 5.5 to about 7.0 of the above pharmaceutical composition, but the above pharmaceutical composition has without  
20        formulation-stability. Thus the present inventors have known that amlodipine maleate may be decomposed due to direct contacting with excipients.

25        The pharmaceutical composition comprises the coated particle of amlodipine maleate, as an active substance for prevent of the direct contact. The pharmaceutical composition of the present invention may contain one or a mixture of above two selected from talc, light silicic anhydride, silicon dioxide, etc. for stability of amlodipine maleate and prevent of electricity and moisture during preparation of the pharmaceutical composition. Further, the pharmaceutical composition may include

pharmaceutical acceptable colors for value of production.

The present invention has efficiencies on formulation-stability regardless size of the coated particle such that the size of the particle is unlimited.

5 In the present invention, may be used coating agent for the coated particle of amlodipine maleate, such as a water-soluble polymer group, a saccharine group and a water-insoluble polymer group that may be broken in 0.01 mol/L of hydrochloric acid solution.

10 The water-soluble polymer group may comprise one or a mixture of above two selected from polyvinylpyrrolidone, a cellulose group (hydroxypropylmethylcellulose, hydroxypropylmethylcellulose phthalate, methylcellulose, sodium carboxymethylcellulose, hydroxyethylcellulose, hydroxypropylcellulose, etc.), pectin, galactomannan, gelatin, a polyethyleneglycol group, polymethacrylate, a cyclodextrin group (betacyclodextrin, methylbetacyclodextrin, hydroxypropylbetacyclodextrin), carbomer, 15 polyvinylalcohol, etc. Preferably, coating agents of OPADRY® (Colorcon Co., Ltd.) and Eudragit® (Rohm Co., Ltd.) are used for ease of coating and rapid dissolution rate.

20 The saccharine group, as the coating agent, may comprise sugar, sorbitol, mannitol, etc., which may be used one or a mixture of above two selected from the saccharines. With the coating agent, may be used a mixture of above one selected from excipients, dilluents, surfactants, lubricants, antifoaming agents, plasticizer for ease of manufacture and obtainment of excellent coating.

25 An object of a method of the present invention is accomplished by coating a particle of amlodipine maleate using fluidized bed granulator. The method comprises flowing a particle of amlodipine maleate using fluidized bed granulator and dispersing the coating agent on the particle of amlodipine maleate to thereby

forming the coated particle of amlodipine maleate.

In flowing of the method, amlodipine maleate may be flowed alone in fluidized bed granulator, and/or one or above two of talc, silicic anhydride, silicon dioxide, etc. may be put into the fluidized bed granulator, for stabilizing of  
5 amlodipine maleate and preventing electricity and moisture, with amlodipine maleate, flowed and coated.

Solid formulations, such as capsules, tablets, powders, granules, etc., including the coated particle of amlodipine maleate are prepared by conventional  
10 methods, and may contain, as occasion demands, above 2 or 1 respectively selected from excipients, sweeteners, flavors, dilluents, lubricants, surfactants, thickening agents, pH control agents, stain, antifoaming agent etc. so as to dose into patients directly.

The solid formulation comprising the coated particle of amlodipine maleate  
15 is prepared by coating the particle of amlodipine maleate using fluidized bed granulator and formulizing as tablets, capsules, powders, etc. such that decomposition reaction of amlodipine maleate is prevented during long storage and reference dissolution rate is maintained, accordingly with excellent formulation-stability.

#### **Advantageous Effects**

The present invention provides a coated particle of amlodipine maleate for preventing a loss of amlodipine maleate during storage and distribution. The present invention also provides a pharmaceutical composition comprising the coated particle  
25 of amlodipine maleate for treating cardiovascular disease(CVD). The pharmaceutical composition has efficiencies on sufficient dissolution rate, stable bioavailability, formulation-stability equal to or more than one of amlodipine

besylate during long storage, and/or formulation-stability enhanced regardless pH.

### **Brief Description of the Drawings**

FIG. 1 represents a graph of blood concentration of formulations comprising  
5 a uncoated particle of amlodipine maleate (AM1) and a coated particle of  
amlodipine maleate(AM-8) prepared in Example 5.

### **Best Mode For Carrying Out the Invention**

The following examples are intended to describe the present invention in  
10 further detail and should not be constructed as limiting the scope of the present  
invention.

#### **<Example 1>**

Amlodipine maleate (Manufacturer: UniChem) 320g and talc 200g was  
15 stirred in fluidized bed granulator, OPADRY-AMB® 700g and talc 300g was  
dispersed in water 10kg and then the dispersed solution was sprayed into the  
granulator to form a coated amount of 170 weight percent over powder of  
amlodipine maleate. The resultant particle was thoroughly dried in the granulator  
and sieved with No. 35 sieve to form coated particles of amlodipine maleate.

20

#### **<Example 2>**

Amlodipine maleate (Manufacturer: Kyongbo Pharm. Co., Ltd) 320g and  
talc 200g was stirred in fluidized bed granulator, OPADRY-AMB® 700g and talc  
300g was dispersed in water 10kg and then dispersed solution was sprayed into the  
25 granulator to form a coated amount of 170 weight percent over powder of  
amlodipine maleate. The resultant particle was thoroughly dried in the granulator  
and sieved with No. 35 sieve to form coated particles of amlodipine maleate.



**<Example 3>**

Amlodipine maleate (Manufacturer: Kyongbo Pharm. Co., Ltd) 320g and talc 200g was stirred in fluidized bed granulator, OPADRY-OYC-7000A<sup>®</sup> 700g and talc 300g was dispersed in methylene chloride 3kg and anhydrous ethanol 7kg and then dispersed solution was sprayed into the granulator to form a coated amount of 170 weight percent over powder of amlodipine maleate. The resultant particle was thoroughly dried in the granulator and sieved with No. 35 sieve to form coated particles of amlodipine maleate.

**<Example 4>**

Amlodipine maleate (Manufacturer: Kyongbo Pharm. Co., Ltd) 320g and talc 200g was stirred in fluidized bed granulator, sugar 700g and talc 300g was dispersed in water 10kg and then dispersed solution was sprayed into the granulator to form a coated amount of 170 weight percent over powder of amlodipine maleate. The resultant particle was thoroughly dried in the granulator and sieved with No. 35 sieve to form coated particles of amlodipine maleate.

**<Example 5>**

Amlodipine maleate (Manufacturer: Kyongbo Pharm. Co., Ltd) 320g and talc 200g was stirred in fluidized bed granulator, OPADRY-AMB<sup>®</sup> 700g and Anhydrous dibasic calcium phosphate 600g was dispersed in water 12kg and then dispersed solution was sprayed into the granulator to form a coated amount of 223 weight percent over powder of amlodipine maleate. The resultant particle was thoroughly dried in the granulator and sieved with No. 35 sieve to form coated particles of amlodipine maleate.

**<Preparation of tablet>**

After mixing the coated particles prepared in the above examples respectively in a ratio of the below table 1, tablets of 200mg per each tablet are  
 5 produced using rotary tablet press machine.

	AM-1	AM-2	AM-3	AM-4	AM-5
Amlodipine maleate (as amlodipine)	6.4g (5.0g)	6.4g (5.0g)	-	-	
Coated particle of example 1 (as amlodipine)	-	-	28.1g (5.0g)	-	-
Coated particle of example 2 (as amlodipine)	-	-	-	28.1g (5.0g)	-
Coated particle of example 3 (as amlodipine)					28.1g (5.0g)
Anhydrous dibasic calcium phosphate	63.0g	63.0g	63.0g	63.0g	63.0g
Microcrystalline cellulose	124.6g	122.6g	100.9g	100.9g	100.9g
Sodium starch glyconate	4.0g	6.0g	6.0g	6.0g	6.0g
Magnesium stearate	2.0g	2.0g	2.0g	2.0g	2.0g
pH	5.7	5.8	6.1	5.9	6.2

	AM-6	AM-7	AM-8	AM-9	AM-10
Amlodipine maleate	-	-	-	6.4g (5.0g)	-
Coated particle of example 3 (as amlodipine)	-	-	-	-	28.1g (5.0g)
Coated particle of example 4 (as amlodipine)	28.1g (5.0g)	-	-	-	-
Coated particle of example 5 (as amlodipine)	-	33.6g (5.0g)	33.6g (5.0g)	-	-
Anhydrous dibasic calcium phosphate	63.0g	63.0g	63.0g	63.0g	63.0g
Microcrystalline cellulose	100.9g	95.4g	92.9g	120.6g	98.9g
Sodium starch glyconate	6.0g	6.0g	6.0g	6.0g	6.0g
Succinic acid	-	-	2.5g	2.0g	2.0g
Magnesium stearate	2.0g	2.0g	2.0g	2.0g	2.0g
pH	6.3	6.0	4.9	4.7	4.8

In the above tables, AM represents amlodipine maleate.

**<Experimental example 1> Test of dissolution rate and content loss for formulation-stability**

An experiment of dissolution rate was carried out by dissolving respectively the above-prepared tablets and a tablet containing amlodipine besylate, has been sold (novasc<sup>®</sup>, containing 5mg as amlodipine), in 500 mL of 0.01mol/L hydrochloric aqueous solution at a temperature of 37°C for 30 minutes at 75 rpm. And an experiment of content loss, with respect to initial content, was performed by storage as opened at a relative humidity (R.H) of 75 weight percent and 100 weight percent respectively for 3 months. The result is represent in the below table.

	AM-1	AM-2	AM-3	AM-4	AM-5
Dissolution rate	89.5%	91.1%	92.6%	91.3%	86.5%
Content loss in a condition of 40°C and R.H. 75% after 3 months	4.6%	13.8%	0.2%	0.1%	1.3%
Content loss in a condition of 50°C and R.H. 100% after 3 months	11.5%	-	5.6%	-	6.1%

	AM-6	AM-7	AM-8	AM-9	AM-10	Novasc <sup>®</sup>
Dissolution rate	84.6%	90.6%	91.0%	91.0%	92.0%	88.5%
Content loss in a condition of 40°C and R.H. 75% after 3 months	1.0%	0.0%	1.0%	14.2%	1.9%	1.1%
Content loss in a condition of 50°C and R.H. 100% after 3 months	-	4.9%	5.8%	35.2%	6.8%	7.0%

In the above table, AM represents amlodipine maleate.

The above table showed that formulation-stability of the tablet comprising the coated particles of amlodipine maleate, compared with one of Novasc<sup>®</sup>, was excellently enhanced regardless pH of tablets. However the tablets (AM-1 and AM-2), containing uncoated particles of amlodipine maleate, had low formulation-stability in a tablet pH range of 5.5 to 7.0 compared with one of Novasc<sup>®</sup>. Further the tablet (AM-9), containing uncoated particles of amlodipine maleate, out of the tablet pH range had rapidly content loss.

Accordingly, the pharmaceutical composition, comprising the coated particle of amlodipine maleate according to the present invention prevents content loss of amlodipine maleate during storage and circulation, and has excellent efficiencies on formulation-stability equal to or more than tablet containing amlodipine besylate and regardless pH.

#### **<Experimental example 2> Bioequivalence test**

To confirm high bioavailability of amlodipine maleate, bioequivalence test was carried out on tablets of the present invention and the conventional tablet.

Into 12 men at an age of 19 to 55 dosed the tablet (AM-1) comprising amlodipine maleate and the tablet (AM-8) containing the coated particle of amlodipine maleate respectively, and after 1, 2, 4, 6, 8, 10, 12, 14, 24, 48, 72, 96, 120 and 144 hours, blood concentration of amlodipine was measured. The result referred to Figure 1. And proper of pharmacokinetics was represented in the below table(Testing laboratory: Asan medical center).

Example	Tablet of AM-1	Tablet fo AM-8
T <sub>max</sub> (hr)*	6.00 (2.00 - 14.00)	8.00 (2.03 - 23.35)
C <sub>max</sub> (ng/mL)	4.74 ± 2.42	4.68 ± 2.62
AUC(0→∞) (ng·hr/mL)	168.09 ± 94.12	158.97 ± 88.38
T <sub>1/2β</sub> (hr)	35.28 ± 17.52	34.79 ± 20.29
MRT (hr)	43.25 ± 7.63	43.55 ± 8.39

\* mean (range)

As showed in the above table, the pharmaceutical composition comprising the coated particle of amlodipine maleate has excellent efficiencies on property of pharmacokinetics equal to or more than conventional compositions containing amlodipine maleate according to testing, such as time that takes to blood maximum concentration of drug (T<sub>max</sub>), blood maximum concentration of drug (C<sub>max</sub>), bioavailability (AUC<sub>0→∞</sub>), half life (T<sub>1/2β</sub>) and staying time of drug in blood.

Therefore the pharmaceutical composition of the present invention has excellent dissolution rate with respect to reference and stable bioavailability equal to or more than the conventional pharmaceutical composition comprising uncoated particles of amlodipine maleate.